AMENDMENTS TO THE CLAIMS:

- 1. (Currently amended) A process for converting a biomass into a blending component for a petroleum-derived fuel comprising:
- (a) extracting a lignin-containing fraction in a reaction medium of water from the biomass to provide a lignin feed material;
- (b) <u>base-catalyzed</u> depolymerizing the lignin feed material in <u>the presence of water an</u> aqueous solvent to provide a first composition comprising a depolymerized lignin; and
- hydrodeoxygenation and hydrocracking first composition to provide a second composition comprising an aromatic hydrocarbon comprising C₇-C₁₀ alkylbenzenes, wherein the second composition provides for use as a blending component for a petroleum or petroleum-derived fuel.
- 2. (Canceled)
- 3. (Canceled)
- 4. (Original) The process of claim 1 wherein the blending composition is further defined as having a blending octane number of about 110 or higher.
- 5. (Canceled) The process of claim 1 wherein the depolymerization is further defined as a base-catalyzed depolymerization.
- 6. (Canceled) The process of claim 2 or 3 wherein the aromatic hydrocarbons are further defined as comprising C_7 to C_{10} alkylbenzenes.
- 7. (Original) The process of claim 1 wherein the biomass is a lignocellulose biomass.
- 8. (Original) The process of claim 1 wherein the second composition is further defined as comprising about 5% to 40% alkylated napthenes.
- 9. (Original) The process of claim 8 wherein the second composition comprises about 5% to 30% alkylated napthenes.
- 10. (Currently Amended) The process of claim 6 $\underline{1}$ wherein the second composition is further defined as comprising from about 75% to about 95% C_7 to C_{10} alkylbenzenes.
- 11. (Original) The process of claim 1 wherein the biomass is further defined as comprising from about 5% to about 70% lignin.
- 12. (Original) The process of claim 11 wherein the biomass comprises about 50% lignin.

- 13. (Original) The process of claim 1 or 12 wherein the biomass is further defined as comprising a Kraft lignin, organosolve lignin, a lignin derived from wood processing, a lignin as an ethanol process by-product, a lignin from a by-product of pulp and paper processing, or a combination thereof.
- 14. (Original) The process of claim 1 wherein the aqueous solvent comprises a dilute alkali hydroxide solution.
- 15. (Original) The process of claim 14 wherein the alkali hydroxide solution is about 0.5% to about 10% wt of an alkali hydroxide.
- 16. (Original) The process of claim 15 wherein the alkali hydroxide solution is about 0.5% to about 3% by wt NaOH.
- 17. (Original) The process of claim 1 wherein the depolymerization is carried out in the presence of a CsX-type zeolite as a superbase catalyst.
- 18. (Original) The process of claim 17 wherein the solid CsX-type zeolite superbase catalyst is used together with a 0.5% 5% alkali hydroxide solution as a co-catalyst system.
- 19. (Currently Amended) The process of claim 5 1 wherein the base-catalyzed depolymerization is further defined as occurring at an operational temperature of from about 300°C to about 340°C.
- 20. (Original) The process of claim 19 wherein the base-catalyzed depolymerization operation temperature is from about 310°C to about 330°C.
- 21. (Currently Amended) The process of claim 5 1 wherein the lignin-containing feedstock material has a liquid hourly space velocity of from 0.5h⁻¹ to about 10h⁻¹.
- 22. (Original) The process of claim 21 wherein the liquid hourly space velocity is from about $2.0h^{-1}$ to about $9.0h^{-1}$.
- 23. (Canceled)
 - 24. (Currently Amended) The process of claim 23 1 wherein hydrodeoxygenation of the first composition depolymerized lignin is further defined as catalyzed by a MMo/γ-Al₂O₃ catalyst and hydrocracking of the first composition depolymerized lignin is further defined as catalyzed by a sulfided MMo/SiO₂-Al₂O₃-zeolite catalyst, wherein M is a Group VI to VIII transition metal promoter.

- 25. (Original) The process of claim 24 wherein the ratio of MMo/ γ Al₂O₃ catalyst: sulfided MMo/SiO₂-Al₂O₃ zeolite catalyst is from 1:1 to 9:1.
- 26. (Cancelled).
- 27. (Original) The process of claim 1 wherein the hydroprocessing is further defined as comprising a hydrogen pressure of from about 500 psig to about 1800 psig.
- 28. (Original) The process of claim 27 wherein the hydrogen pressure is from about 500psig to about 1000 psig.
- 29. (Original) The process of claim 1 wherein the hydroprocessing is further defined as comprising a reaction temperature of from about 360°C to about 390°C.
- 30. (Original) The process of claim 29 wherein the reaction temperature is from about 380°C to about 390°C.
- 31. (Original) The process of claim 24 wherein M is selected from the group consisting essentially of Ru, Co, Re, Cr, Fe, Pt, and combinations thereof.
- 32. (Original) The process of claim 1 wherein the lignin feed material is depolymerized in the absence of alcohol.
- 33. (Withdrawn) A biomass-derived blending component for a petroleum or petroleum-derived fuel comprising about 70% to about 95% C_7 to C_{10} alkylbenzene.
- 34. (Withdrawn) The biomass-derived blending component of claim 33 further defined as comprising about 5% to about 30% alkylated naphthenes.
- 35. (Withdrawn) The biomass-derived blending component of claim 33 further defined as comprising about 5% to about 10% alkylated napthenes.
- 36. (Withdrawn) A method for enhancing the octane number of a petroleum or petroleum-derived fuel comprising: combining the biomass derived blending component of claim 33 with a petroleum or petroleum-derived fuel at a ratio of about 1:10 to about 1:4 by volume to provide a blended fuel, wherein the octane level of the petroleum blended fuel is enhanced about 1% to about 30% over the octane level of the petroleum or petroleum-derived based fuel without the blending component.
- 37. (Withdrawn) The method of claim 36 wherein the octane level of the blended fuel is enhanced about 30% over the petroleum or petroleum-derived fuel without the blending component.

- 38. (Withdrawn) The method of claim 36 wherein the petroleum or petroleum-derived fuel is gasoline.
- 39. (Currently Amended) A process for converting a biomass into a blending component comprising a monocyclic aromatic hydrocarbon-rich composition for petroleum-derived fuels comprising:
- a) dispersing a lignin-containing feedstock material in a reaction medium of water to provide a dispersed lignin composition;
- b) <u>base-catalyzed</u> depolymerizing the dispersed lignin composition in the presence of water to provide a depolymerized lignin product; and
- c) hydroprocessing the depolymerized lignin product <u>by sequential or simultaneous</u> <u>hydrodeoxygenation and hydrocracking</u> to produce a blending component comprising <u>monocyclic aromatic hydrocarbons</u> <u>C₇ to C₁₀ alkylbenzene</u>.
- 40. (Canceled).
- 41. (Original) The process of claim 39 wherein the blending component comprises a blending octane number of about 95-150.
- 42. (Original) The process of claim 39 wherein the blending component is further defined as having a blending octane number of about 110.
- 43. (Canceled).
- 44 (Currently Amended) A process for producing BTX (benzene, toluene and xylenes) aromatic comprising:
 - a) extracting lignin-containing material from a biomass;
 - b) dispersing the lignin-containing material in a reaction medium of water;
- c) subjecting the dispersed lignin-containing material to an alkali hydroxide solvent in the presence of water to produce a first composition comprising a depolymerized lignin; and
- d) hydroprocessing the <u>depolymerized lignin by sequential or simultaneous</u>

 <u>hydrodeoxygenation and hydrocracking first composition</u> to provide a second composition

 comprising benzene, toluene, and xylenes, wherein the benzene, toluene, and xylenes are <u>used as</u>

 intermediates in the production of other organic chemicals.
- 45. (Currently Amended) A process for converting a biomass into a blending composition for a petroleum-derived fuel comprising:

- a) extracting lignin-containing material from a biomass;
- b) dispersing the lignin-containing material in a reaction medium of water;
- c) subjecting the dispersed lignin-containing material to a alkali hydroxide solution to produce a first composition comprising a depolymerized lignin;
- d) hydrodeoxygenating and hydrocracking the first composition to produce a second composition comprising C_7 to C_{10} alkylbenzenes, wherein the second composition provides is used as a blending component for a petroleum-derived fuel.
- 46. (Original) The process of claim 45 wherein the hydrodeoxygenating and hydrocracking of the first composition occur substantially simultaneously.
- 47 (Original) The process of claim 45 wherein the alkali hydroxide solution is about 1% to about 10% wt of an alkali hydroxide.
- 48. (Original) The process of claim 47 wherein the alkali hydroxide solution is about 1% to about 3% by wt NaOH.
- 49. (Canceled).
- 50. (Original) The process of claim 47 wherein the second composition is further defined as comprising from about 75% to about 95% C_7 to C_{10} alkylbenzenes.